

VITAMIN K AND ESSENTIAL FATTY ACIDS

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Vitamin K is a general name for a group of compounds all with similar biological activity. They all contain the 2-methyl-1,4-naphthoquinone nucleus with a lipophilic side chain at position 3. The three best known members are phyloquinone (vitamin K1) which is the only type of vitamin K found in plants. Vitamin K2, the menaquinones, consists of a family of compounds with variable length isoprenyl side chains. Vitamin K3, menadione, is a pro-vitamin which can be converted to vitamin K2 by animals. Menadione and the menaquinones may occasionally have toxic effects in high doses whereas phyloquinone seems to be safe even in massive overdose. Phyloquinone is therefore the preferred form of the vitamin for human use.

Vitamin K compounds are widely distributed in foods. Among animal foods, eggs, butter and liver are good sources and contain amounts of from about 2 to about 50 $\mu\text{g}/100\text{g}$ of the food. Green vegetables are also good sources and may contain from 30 to as much as 800 $\mu\text{g}/100\text{g}$ of the food. Spinach, kale, sprouts and broccoli are good sources. Vegetable oils, and products made from vegetable oils such as margarines and salad dressings, can also be good sources, containing from 10 to 300 μg of vitamin K per 100g of oil. Olive oil and soy oil are particularly rich in vitamin K. Some vitamin K is also made from gut bacteria although this is difficult to quantitate and may be very variable.

The US Recommended Daily Allowance (RDA) for vitamin K starts at 10 $\mu\text{g}/\text{day}$ for infants and rises to

65 $\mu\text{g/day}$ in women and 80 $\mu\text{g/day}$ in men. There is, however, evidence that vitamin K from some foods may be relatively poorly absorbed and there have been suggestions that these RDAs for ordinary foods may be somewhat low (BLMG Gijsbers et al, Effect of food composition on vitamin K absorption in human volunteers, Br J Nutrition 1996; 76: 223-229).

The best known role for vitamin K in humans is as a co-factor for the synthesis of six of the proteins involved in blood clotting. These proteins are inactive proenzymes which are converted to active enzymes in the presence of calcium during the coagulation process. These proteins contain an unusual amino acid, gamma-carboxy-glutamate. This is formed by the carboxylation of glutamic acid residues in the protein by the enzyme gamma-glutamyl carboxylase, in a vitamin-K dependent reaction. In the absence of vitamin K, the normal forms of the clotting factors cannot be synthesised. Proteins containing gamma-carboxy-glutamate have become known by the general name of Gla proteins.

For some time it was thought that Gla proteins were confined to the clotting system and it was largely on this basis that the RDAs were estimated. However, it is now known that enzymes with gamma-glutamyl-carboxylase activity are widely distributed in many different tissues and so it is probable that there are many functions of Gla proteins to be discovered. These proteins are particularly abundant in kidney and in bone and so it is assumed that they have particular roles to play in these organs. Two Gla proteins are particularly abundant in bone. Bone Gla

protein (BGP, commonly known as osteocalcin) contains three Gla residues and is present in great abundance in bone, dentin and cartilage. Matrix Gla protein (MGP) is also found in substantial amounts in bone,
5 dentin and cartilage. Much ongoing research is trying to identify the roles of these proteins which seem to be involved in determining the strength and resilience of the structure. The kidney Gla proteins may be involved in regulation of calcium excretion so
10 that vitamin K may play a role in integrating the various mechanisms involved in maintaining bone strength (NC Binkley and J W Suttie, Vitamin K nutrition and osteoporosis, J Nutr 1995; 125: 1812-21 and C Vermeer et al, Effects of vitamin K on bone
15 mass and bone metabolism, J Nutr 1996; 126: 1187S-1191S).

Recently there is evidence that vitamin K can have clinically relevant effects on bone. In women with osteoporosis, a controlled study showed that 45mg/day
20 of vitamin K2 could reduce the risk of bone fractures and slow down but not prevent a progressive loss of bone mineral density (M Shiraki et al, J Bone Mineral Res 2000; 15: 515-521). In a prospective study of
25 72,000 nurses, women with the lowest quintile of vitamin K intake (109 μ g/day and below) had an increased risk of fractures D Feskanich et al, Vitamin K intake and hip fractures in women: a prospective study, Am J Clin Nutr 1999; 69: 74-79). In an older group of men and women, mostly over 70,
30 there was a progressively reducing risk of osteoporotic fracture as vitamin K intake increased. The lowest risk was in the highest quartile of vitamin K intake of more than 262 μ g/day in women and

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more than 234 $\mu\text{g/day}$ in men (SL Booth et al, Dietary vitamin K intakes are associated with hip fracture but not with bone mineral density in elderly men and women, Am J Clin Nutr 2000; 71: 1201-8). The
5 progressive effect of increasing daily intakes suggests that doses of vitamin K much higher than the RDAs may be important in bone health, especially in older people.

The essential fatty acids (EFAs) are a completely
10 different group of nutrients. There are two types, the omega-6 derived from the parent linoleic acid, and the omega-3 derived from the parent alpha-linolenic acid (figure 1). The EFAs cannot be synthesised de novo by humans or other mammals. Nor
15 can mammals convert omega-3 EFAs into omega-6 EFAs or vice versa. Mammals can interconvert one omega-6 EFA into another omega-6 EFA via the pathways shown in figure 1. Similarly, omega-3 EFAs can be converted one to another. The pathways shown in figure 1
20 usually progress from linoleic acid or alpha-linolenic acid downwards, but retro-conversions are possible.

The EFAs are essential components of complex lipids such as triglycerides, cholesterol esters and
25 phospholipids, and are absolutely required for the normal structure and function of all cell and other membranes in the body. Deficiencies of EFAs cause widespread defects in all body systems. While dietary deficiencies of the parent EFAs, linoleic and alpha-linolenic acids are relatively rare, deficiencies of
30 their metabolites are relatively common because of impaired conversion mechanisms. As a result low

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levels of fatty acids like dihomogammalinolenic acid (DGLA), arachidonic acid (AA) and adrenic acid (AdrA) of the n-6 series and of stearidonic acid (SA), eicosapentaenoic acid (EPA), docosapentaenoic acid (DPA) and docosaheptaenoic acid (DHA) of the n-3 series have been commonly reported. Such low levels have been found in skin diseases including atopic eczema; reproductive system disorders including premenstrual syndrome, breast pain and menstrual pain; diabetes; cardiovascular disorders; bone disorders; kidney diseases; psychiatric diseases including schizophrenia, depression, stress and attention deficit hyperactivity disorder; and many other conditions. Treatment with EFAs, especially with gamma-linolenic acid (GLA) of the n-6 series and with EPA and DHA of the n-3 series has been reported to be associated with a wide range of beneficial effects. These effects have been reported to be enhanced by certain nutrients such as zinc and vitamin B6 which are important in EFA metabolism.

The present invention is based on the inventor's finding of a completely unexpected and hitherto unreported interaction between vitamin K and EFAs.

The present invention provides nutritional and pharmaceutical formulations comprising in combination a source of vitamin K and a source of at least one essential fatty acid (EFA), in which the concentration of vitamin K is not less than 1000 $\mu\text{g}/100\text{g}$. Preferably, the concentration of vitamin K is not less than 1000 $\mu\text{g}/10\text{g}$. The formulations of the invention preferably comprise between 50 μg and 100 mg vitamin K and between 50 mg and 100 g of the

EFA. These are to provide a daily dose of these amounts and the formulation may be in the form of a single dosage, to provide these intakes in one go, or in the form of divided doses.

5 The present invention further provides nutritional and pharmaceutical formulations comprising in combination a source of Vitamin K and a source of at least one EFA, but which exclude proteins or amino acids as part of the active ingredients of the
10 formulations.

Vitamin K is preferably in the form of phylloquinone (vitamin K1).

The EFA may be selected from the n-6 series: gamma-linolenic acid, dihomogammalinolenic acid,
15 arachidonic acid and adrenic acid, and combinations of these EFAs. Alternatively, the EFA is selected from the n-3 series: stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexaenoic acid, and combinations of these EFAs.
20 In a further embodiment of the present invention at least one n-6 EFA is present with at least one n-3 EFA, each EFA selected from the above lists.

The active ingredient of the nutritional or pharmaceutical composition may consist essentially
25 wholly of EFA and vitamin K or, alternatively, the formulations of the present invention may further comprise one or more essential vitamins and/or minerals or one or more pharmaceutical drugs.

The present invention further provides foodstuff which already contain EFAs to which have been added vitamin K in an amount to raise the vitamin K content of the food to 1000 μg / 100 g food, or more. The
5 specific EFA(s) content may also be raised artificially by the addition of one of more EFAs.

It is desirable that the present invention still further provides foodstuffs containing EFAS but excluding proteins or amino acids as part of the
10 active ingredients. Added to the composition is vitamin K to artificially raise the vitamin K content of the foodstuff. The specific EFA(s) content may also be raised artificially by the addition of one of more EFAs.

15 The present invention still further provides foodstuff which naturally contains clinically or nutritionally small amounts of vitamin K and / or EFA(s) but to which has been added vitamin K and EFAs, for example to the dosage regime of the
20 formulations of the first aspect of the present invention. Milk and other dairy products or simulated dairy products are particularly appropriate for this type of enrichment.

The foodstuffs and nutritional or pharmaceutical
25 formulations of the present invention may be used to treat or prevent a variety of diseases or conditions. These may include:

premenstrual or menstrual disorders of any kind;
bone or calcium disorders of any kind, including
30 osteoporosis;

metabolic or cardiovascular disorders including diabetes, obesity, elevated blood cholesterol or triglyceride levels or cardiovascular disorders;

5 stress, mental, psychological, psychiatric or neurological disorders;

skin disorders;

asthma or other respiratory disorder;

10 arthritis or any form of inflammatory, gastrointestinal, kidney or reproductive system disorder.

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The present invention further provides a method of treatment or prevention of diseases or conditions, including those mentioned above, by the administration of a combination of vitamin K and an
15 EFA, preferably at the dosage rate of between 50 μ g and 100 mg vitamin K and between 50 mg and 100 g EPA. In particular, the disorders to be treated are skin disorders and premenstrual or menstrual conditions. Bone disorders are also of particular importance.

20 The vitamin K may be provided in any form which has biological vitamin K activity in mammals. However, because of its safety and known activity, vitamin K1 (phylloquinone) is the preferred form. The formulations may provide for an increase in vitamin K
25 intake in a nutritional or pharmaceutical formulation or food of from 50 μ g to 100 mg per day. At the same time the formulations should provide for an increase in the intake of one or more of the desired EFAs of between 50mg and 50g per day. Depending on the
30 problem to be addressed, any of the EFAs shown in figure 1 may be used. Linoleic acid, alpha-linolenic acid, GLA, DGLA, AA, AdrA, SA, EPA, DPA and DHA are

likely to be preferred ingredients for particular purposes. The EFAs may be provided as purified or partially purified compounds or may be supplied by natural oils which are rich in one or more EFAs. For example, borage and evening primrose oils are good sources of GLA, Echium oils are good sources of SA, marine oils are often good sources of EPA, DPA, DHA and sometimes AA, while oils from various microbial sources, including fungal and algal oils can be sources of GLA, DGLA, AA, SA, EPA or DHA. The EFAs can be in any chemical form which is absorbed into the body and incorporated into body lipids. Such forms include but are not limited to free acids, sodium, potassium, lithium and other salts, triglycerides and other glycerides, cholesterol, ethyl, methyl and other esters, amides, and phospholipids.

The vitamin K and the EFA when used for pharmaceuticals or nutritional supplements can be incorporated into any appropriate dosage form known to those skilled in the art. Such dosage forms include soft and hard gelatin capsules, tablets, microcapsules of various types, powders and carriers of various types, liquids, emulsions, micelles and any other forms. Flavourants, colourants, emulsifiers and conventional diluents and excipients may be included, alone or in combination. Examples of formulations of the dosage follow.

Example 1

500 mg or 100 mg hard or soft gelatin capsules in which a natural oil containing GLA is formulated with

vitamin K, preferably phylloquinone, at a level between 0.05 and 1.0 mg per capsule.

Example 2

500 mg or 1000 mg hard or soft gelatin capsules where
5 the natural oil contains stearidonic acid (SA),
eicosapentaenoic acid (EPA), docosapentaenoic acid
(DPA), docosahexaenoic acid (DHA),
dihomogammalinolenic acid (DGLA) or arachidonic acid
(AA).

10 Example 3

500 mg or 1000 mg hard or soft gelatin capsules
containing gamma-linolenic acid (GLA) in either
triglyceride or ethyl-ester form in which the purity
of the GLA is greater than 50% and preferably greater
15 than 90% and in which vitamin K, preferably
phylloquinone, is provided at a level of between 0.05
and 5.0 mg/capsule.

Example 4

500 mg or 1000 mg hard or soft gelatin capsules
20 containing SA, EPA, DPA, DHA, DGLA or AA or a mixture
of these fatty acids and in which vitamin K,
preferably phylloquinone, is provided at a level of
between 0.05 and 5.0 mg/capsule.

Example 5

25 Liquid natural oils containing:
GLA; or

SA, EPA, DPA, DHA, DGLA or AA; or
GLA in triglyceride or ethyl-ester form in which
the purity of the GLA is greater than 50% and
preferably greater than 90%;

5 or to which are added 1000 micrograms/100g and
100mg/100g of vitamin K, preferably in the
phyllloquinone form. Such oils may be used
themselves, or flavoured using appropriate
flavourings, or incorporated into microcapsules made
10 from any appropriate material or added to foodstuffs
of any appropriate type.

Example 6

Milk or milk products from any edible source, either
animal or vegetable, such as soy milk, to which are
15 added phylloquinone to raise the concentration to
over 1000 microg/100 g and preferably to over 5000
microg/100 g together with one or more fatty acids
selected from GLA, DGLA, AA, SA, EPA, DPA and DHA, to
raise the concentration of each selected fatty acid
20 to more than 100 mg/100 g and preferably to more than
1000 mg/100 g.

When used in foods, the formulations may be prepared
by increasing the concentration of vitamin K in the
food to 1000 μ g/100g or more. With some foods, such
25 as milks, dairy products or vegetable oils, moderate
amounts of EFAs may already be present in the natural
food. Increasing the vitamin K content of such foods
to a level above that present in any natural food is
within the framework of the invention.

Alternatively, in addition to raising the vitamin K content of an EFA-containing food to over 1000 $\mu\text{g}/100\text{g}$, the desired EFA may also be added to the food to raise the amount provided. Natural and soy or other vegetable-based milks, soy and related products, dairy products including yogurts, cheeses, butters, margarines, or any other types of foods may all be treated in this way to provide a combination of vitamin K and an EFA.

These formulations may be used for general health purposes, or for specific conditions where either EFAs or both have been found to be helpful. These conditions include premenstrual and menstrual disorders, skin disorders, diabetes, elevated cholesterol and triglyceride levels, cardiovascular disorders, arthritis and any form of inflammatory disorder, respiratory disorders such as asthma, gastro-intestinal, urinary and reproductive system disorders in both sexes, mental, psychological and psychiatric disorders such as stress, chronic fatigue, behavioural problems in children and adults, depression, alcoholism and more serious psychiatric disorders such as schizophrenia and bipolar disorder, neurological disorders such as Parkinsonism, and any form of dementia and any other form of illness in which the combinations are found to be helpful. Osteoporosis and related disorders of bone and calcium metabolism are likely to provide particularly important uses for the invention.

Brief Description of the Figures

Fig. 1 The n-6 and n-3 essential fatty acids

Experimental Data

A woman with atopic dermatitis and with mild premenstrual syndrome was recommended to take 3g/day of evening primrose oil (EPO). EPO is a widely used nutritional supplement for these problems. It contains about 70% of linoleic acid, but more importantly contains 8-12% of GLA which can by-pass a block in the conversion of linoleic acid to GLA which can occur in many situations, including atopic dermatitis, stress and menstrual disorders. Not everyone responds to EPO but this is an exceptionally safe nutritional supplement and does not cause any important side effects. However, to my surprise in this woman the EPO not only failed to have any therapeutic effect but caused a range of unusual side effects including facial reddening and rashes, a worsening of her dermatitis, gastro-intestinal disturbances and anxiety and depression. As a result of a series of investigations, she was found to have a vitamin K deficiency, possibly partly due to dietary problems and partly due to gastrointestinal infections which had necessitated the use of antibiotics which had probably changed her gut bacteria. The vitamin K deficiency was corrected by vitamin K1 supplements but this did not improve her skin or her premenstrual syndrome. As an experiment she was then cautiously given EPO again. This time there were no adverse effects at all, her skin improved and her premenstrual syndrome was resolved. This case suggested a hitherto unsuspected and important positive interaction between vitamin K and EFAs.

A second woman presented with severe menstrual cramps and mild premenstrual syndrome. I suggested that she should take a low dose of EPO (1g/day) to help with her premenstrual syndrome and a higher dose (4g/day) of a fish oil containing 23% of EPA and 8% of DHA to help with her menstrual cramps. Unfortunately she showed no response in either of her problems. She had a normal diet and no evidence of vitamin K deficiency but I wondered whether giving vitamin K might help. She therefore took 2mg (2000 μ g) per day of a vitamin K1 supplement for a month which also had no effect on her menstrual problems. However, on reintroducing the EPO and fish oil, her premenstrual syndrome disappeared completely and her menstrual pain was greatly reduced.

These cases show that vitamin K can greatly improve the therapeutic effects of EFAs, reducing any side effects and enhancing therapeutic effects.

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